Application No.: 10/828,795 Filing Date.: April 21, 2004

AMENDMENTS TO THE CLAIMS

1-49. (Canceled).

- 50. (Currently amended) A composition for affecting weight loss comprising-a weight loss affecting amount of a combination of:
 - (a) naltrexone or a pharmaceutically acceptable salt thereof; and
- (b) a sustained release formulation of bupropion or a pharmaceutically acceptable salt thereof in an amount effective to induce weight loss in an individual; and
- (b) naltrexone or a pharmaceutically acceptable salt thereof in an amount effective to enhance the weight loss effect of the bupropion or salt thereof;

wherein said composition is formulated for oral administration in a single oral dosage form.

- 51. (Currently amended) The composition of claim 50, wherein said combination composition comprises about 5 mg to about 50 mg of naltrexone or a pharmaceutically acceptable salt thereof.
- 52. (Currently amended) The composition of claim 50, wherein said combination composition comprises about 30 mg to about 300 mg of bupropion or a pharmaceutically acceptable salt thereof.
- 53. (Currently amended) The composition of claim 50, wherein said eombination composition comprises about 5 mg to about 50 mg of naltrexone or a pharmaceutically acceptable salt thereof, and about 30 mg to about 300 mg of bupropion or a pharmaceutically acceptable salt thereof.
- 54. (Withdrawn) The composition of claim 50, further comprising zonisamide or a pharmaceutically acceptable salt thereof.
- 55. (Currently amended) A pharmaceutical composition for affecting weight loss comprising a weight loss affecting amount of a combination of:
 - (a) naltrexone or a pharmaceutically acceptable salt thereof; and
- (b) a sustained release formulation of bupropion or a pharmaceutically acceptable salt thereof in an amount effective to induce weight loss in an individual:
- (b) naltrexone or a pharmaceutically acceptable salt thereof in an amount effective to enhance the weight-reducing effect of the bupropion or salt thereof; and

Application No.: 10/828,795 Filing Date.: April 21, 2004

a pharmaceutically acceptable excipient, diluent, or carrier, wherein said composition is formulated into a single oral dosage formfor oral administration.

- 56. (Currently amended) The pharmaceutical composition of claim 55, wherein said eombination composition comprises about 5 mg to about 50 mg of naltrexone or a pharmaceutically acceptable salt thereof.
- 57. (Currently amended) The pharmaceutical composition of claim 55, wherein said eombination composition comprises about 30 mg to about 300 mg of bupropion or a pharmaceutically acceptable salt thereof.
- 58. (Currently amended) The pharmaceutical composition of claim 55, wherein said eombination composition comprises about 5 mg to about 50 mg of naltrexone or a pharmaceutically acceptable salt thereof, and about 30 mg to about 300 mg of bupropion or a pharmaceutically acceptable salt thereof.
- 59. (Withdrawn) The composition of claim 55, further comprising zonisamide or a pharmaceutically acceptable salt thereof.

60-66. (Canceled).